Initial Approval Date: January 8, 2020

Revised Dates: January 20, 2021; October 14, 2020

### **CRITERIA FOR PRIOR AUTHORIZATION**

Narcolepsy Agents

**BILLING CODE TYPE** For drug coverage and provider type information, see the KMAP Reference Codes webpage.

MANUAL GUIDELINES: Prior authorization will be required for all current and future dose forms available. All

medication-specific criteria, including drug-specific indication, age, and dose for each agent is

defined in Table 1 below.

Oxybate Salts (Xywav™) Pitolisant (Wakix®)

Sodium Oxybate (Xyrem®) Solriamfetol (Sunosi®)

## **GENERAL CRITERIA FOR INITIAL PRIOR AUTHORIZATION:** (must meet all of the following)

- Must be approved for the indication, age, and not exceed dosing limits listed in Table 1.
- For all agents listed, the preferred PDL drug, if applicable, which treats the PA indication, is required unless the patient meets the non-preferred PDL PA criteria.
- Diagnosis of narcolepsy has been confirmed in accordance with International Classification of Sleep Disorders Third Edition (ICSD-3) listed in Table 2.<sup>1</sup>
- If the requested drug is for the treatment of excessive daytime sleepiness (EDS) (must meet both of the following):
  - Patient must have failed a 6-week trial of or have a contraindication to modafinil and/or armodafinil.<sup>2</sup>
  - o Prescriber must provide the patient's baseline Epworth Sleepiness Scale (ESS) score.<sup>3</sup>
- If the requested drug is for the treatment of cataplexy, prescriber must provide patient's baseline frequency of cataplexy episodes per month.
- If the requested drug is for Xywav<sup>™</sup>, prescriber must provide compelling rationale of why the patient will benefit from Xywav<sup>™</sup> over Xyrem<sup>®</sup>.

# **LENGTH OF APPROVAL (INITIAL):** 3 months

### **CRITERIA FOR RENEWAL PRIOR AUTHORIZATION:** (must meet all of the following)

- If the requested drug is for the treatment of EDS, patient has an improvement (reduction) in ESS score.
- If the requested drug is for the treatment of cataplexy, patient has a decrease or maintained a decrease in the number of cataplexy episodes per month.
- Must not exceed dosing limits listed in Table 1.

### **LENGTH OF APPROVAL (RENEWAL): 12 months**

FOR DRUGS THAT HAVE A CURRENT PA REQUIREMENT, BUT NOT FOR THE NEWLY APPROVED INDICATIONS, FOR OTHER FDA-APPROVED INDICATIONS, AND FOR CHANGES TO AGE REQUIREMENTS NOT LISTED WITHIN THE PA CRITERIA:

• THE PA REQUEST WILL BE REVIEWED BASED UPON THE FOLLOWING PACKAGE INSERT INFORMATION: INDICATION, AGE, DOSE, AND ANY PRE-REQUISITE TREATMENT REQUIREMENTS FOR THAT INDICATION.

**LENGTH OF APPROVAL (INITIAL AND RENEWAL): 12 months** 

Table 1. FDA-approved age and dosing limits for Narcolepsy Agents.<sup>4-7</sup>

Agents	Indication(s)	Age	Dosing Limits
	Anale	ptics	
Pitolisant (Wakix®)	Narcolepsy with associated excessive daytime sleepiness  Narcolepsy with cataplexy (type 1 narcolepsy)	≥ 18 years	35.6 mg orally daily CYP2D6 poor metabolizers: 17.8 mg orally daily
Solriamfetol (Sunosi®)	Narcolepsy with associated excessive daytime sleepiness	≥ 18 years	150 mg orally daily
	Psychotherap	eutic Agents	
Oxybate Salts (Xywav™)	Excessive daytime sleepiness associated with narcolepsy  Narcolepsy with cataplexy (type 1 narcolepsy)	≥ 7 years	Adults: 9 grams per night  Pediatrics*:  ≥ 45 kg: 4.5 grams per dose and 9 grams per night 30 to < 45 kg: 3.75 grams per dose and 7.5 grams per night 20 to < 30 kg 3 grams per dose and 6 grams per night
Sodium Oxybate (Xyrem®)	Narcolepsy with associated excessive daytime sleepiness  Narcolepsy with cataplexy (type 1 narcolepsy)	≥ 7 years	Adults: 9 grams per night  Pediatrics*:  ≥ 45 kg: 4.5 grams per dose and 9 grams per night 30 to < 45 kg: 3.75 grams per dose and 7.5 grams per night 20 to < 30 kg 3 grams per dose and 6 grams per night

<sup>\*</sup>There is no specific dosing provided in the manufacturer's labeling (insufficient information) for those weighing under 20 kg. Consider lower initial dosage, lower maximum weekly dosage increases, and lower total maximum nightly dosage.

Table 2. ICSD-3 diagnostic criteria for narcolepsy type 1 and 2.1

Type 1 narcolepsy	Type 2 narcolepsy	
Alternate Names: Hypocretin deficiency syndrome,	Alternate Names: Narcolepsy without cataplexy.	
narcolepsy-cataplexy, narcolepsy with cataplexy.	Diagnostic Criteria: A-E must be met.	
Diagnostic Criteria: A and B must be met.	A. The patient has daily periods of irrepressible	
A. The patient has daily periods of irrepressible	need to sleep or daytime lapses into sleep	
need to sleep or daytime lapses into sleep	occurring for at least three months.	
occurring for at least three months.	B. A mean sleep latency of ≤ 8 minutes and two or	
B. The presence of one or both of the following:	more sleep onset REM periods (SOREMPs) are	
1. Cataplexy (as defined under Essential Features)	found on a MSLT performed according to	
and a mean sleep latency of ≤ 8 minutes and two	standard techniques. A SOREMP (within 15	
or more sleep onset REM periods (SOREMPs) on	minutes of sleep onset) on the preceding	
an MSLT performed according to standard	nocturnal polysomnogram may replace one of	
techniques. A SOREMP (within 15 minutes of	the SOREMPs on the MSLT.	

#### APPROVED PA Criteria

Type 1 narcolepsy	Type 2 narcolepsy	
sleep onset) on the preceding nocturnal	C. Cataplexy is absent.	
polysomnogram may replace one of the	D. Either CSF hypocretin-1 concentration has not	
SOREMPs on the MSLT.	been measured or CSF hypocretin-1	
2. CSF hypocretin-1 concentration, measured by	concentration measured by immunoreactivity is	
immunoreactivity, is either ≤ 110 pg/mL or <1/3	either > 110 pg/mL or > 1/3 of mean values	
of mean values obtained in normal	obtained in normal subjects with the same	
subjects with the same standardized assay.	standardized assay.	
	E. The hypersomnolence and/or MSLT findings are	
	not better explained by other causes such as	
	insufficient sleep, obstructive sleep apnea,	
	delayed sleep phase disorder, or the effect of	
	medication or substances or their withdrawal.	

# References

- 1. Quality measures for the care of patients with narcolepsy. J Clin Sleep Med 2015;11(3):33-355. Available at <a href="http://jcsm.aasm.org/viewAbstract.aspx?pid=29931">http://jcsm.aasm.org/viewAbstract.aspx?pid=29931</a>. Accessed 08/26/2020.
- 2. Practice parameters for the treatment of narcolepsy and other hypersomnias of central origin. *Sleep* 30.12 (2007): 1705-1711. Available at <a href="https://academic.oup.com/sleep/article/30/12/1705/3741350">https://academic.oup.com/sleep/article/30/12/1705/3741350</a>. Accessed on 08/26/2020.
- 3. Johns, Murray W. A new method for measuring daytime sleepiness: the Epworth sleepiness scale. *Sleep* 14.6 (1991):540-545. Available at https://academic.oup.com/sleep/article/14/6/540/2742871. Accessed 12/30/19.
- 4. Wakix (pitolisant) [prescribing information]. Plymouth Meeting, PA: Harmony Biosciences LLC; October 2020.
- 5. Xyrem (sodium oxybate) [prescribing information]. Palo Alto, CA: Jazz Pharmaceuticals; July 2020.
- 6. Sunosi (solriamfetol) [prescribing information]. Palo Alto, CA: Jazz Pharmaceuticals Inc; June 2019.
- 7. Xywav (oxybate salts) [prescribing information]. Palo Alto, CA: Jazz Pharmaceuticals; July 2020.

DRUG UTILIZATION REVIEW COMMITTEE CHAIR	PHARMACY PROGRAM MANAGER	
	DIVISION OF HEALTH CARE FINANCE	
	KANSAS DEPARTMENT OF HEALTH AND ENVIRONMENT	
DATE	DATE	